## THE CLAIMS

## What is claimed is:

- 5 1. A medicated device comprising:
  - a scaffold member suitable for implantation at a tumor or other lesion site; a polymeric coating ("med coat") on the scaffold member; and at least one therapeutic agent in the med coat at a loading sufficient to provide

therapeutic quantities of the therapeutic agent to the site for an extended period of time.

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- 2. The device of claim 1 comprising an anti-cancer therapeutic agent in the med coat.
- 3. The device of claim 1 comprising at least 5 micrograms ( $\mu$ g) of at least one therapeutic agent per square centimeter of the med coat.
- 4. The device of claim 3 comprising at least 50  $\mu g$  of at least one therapeutic agent per square centimeter of the med coat.
- 5. The device of claim 4 comprising at least  $100~\mu g$  of at least one therapeutic agent per square centimeter of the med coat.
- 6. The device of claim 5 comprising at least 500  $\mu$ g of at least one therapeutic agent per square centimeter of the med coat.
- 7. The device of claim 1 comprising sufficient quantity of a therapeutic agent to deliver therapeutically effective quantity of a therapeutic agent into tissue in a region of at least one centimeter from the device.
- 8. The device of claim 7 comprising sufficient quantity of a therapeutic agent to deliver therapeutically effective quantity of a therapeutic agent into tissue in a region of at least two centimeters from the device.

- 9. The device of any one of claims 1-5 wherein the med coat comprises a hybrid polymeric coating comprising a hydrophilic polymer component and a hydrophobic polymer component.
- 5 10. The device of claim 9 wherein the hydrophobic polymer component comprises one or more cellulose ester polymers.
  - 11. The device of claim 1 wherein the polymeric coating comprises an acrylate polymer and PVP/VA copolymer in a weight ratio in the range of from 1.5:1 to 7:1.

12. A medicated device comprising:

a substrate suitable for implantation in a patient's body;

a polymeric coating ("med coat") on the scaffold member; and

at least one therapeutic agent in the med coat at a loading sufficient to provide therapeutic quantities of the therapeutic agent to the patient's tissue in a region in the body extending at least one centimeter from the device.

- 13. The device of claim 12 comprising sufficient quantity of a therapeutic agent to deliver therapeutically effective quantity of a therapeutic agent into tissue in a region of at least two centimeters from the device.
- 14. The device of claim 12 or claim 13 comprising a hybrid polymeric coating comprising a major proportion of one or more hydrophilic polymer materials and a minor proportion of one or more cellulose ester polymers.
- 15. The device of claim 14 wherein the cellulose ester polymer comprises about 3% by weight of the combined weights of the cellulose ester polymer and the hydrophilic polymer materials.
- 30 16. The device of claim 14 wherein the cellulose ester polymer comprises nitrocellulose.

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17. The device of claim 16 wherein the nitrocellulose comprises about 3% by weight of the combined weights of nitrocellulose and the hydrophilic polymer materials.